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LHRH analogs.

The present invention deals with LHRH analogues which contain cytotoxic moleties and have influence on the release of gonadotropins from the pituitary gland of mammals, including humans. The compounds of this invention are represented by the formula:

 $X-R^1-R^2-R^3-Ser-R^5-R^6(Q)-Leu-Arg-Pro-R^{10}-NH_2$ 

wherein

R1 is pGlu, Pro, D-Nal(2), or D-Phe(4Cl),

R<sup>2</sup> is His or D-Phe(4Cl),

R3 is Trp, D-Trp or D-Pal(3),

R5 is Tyr or Arg,

R6 is D-Phe or R6, where R6 is D-Om, D-Lys or D-Phe(NH2),

R10 is Gly or D-Ala,

X is hydrogen, a lower alkanovl group of 2-5 carbon atoms or carbamyl,

Q is bis-(2-chloroethyl)amino group provided that R<sup>5</sup> is D-Phe,

where R<sup>6</sup> is R\*6,

Q is a complexed metal-containing acyl group having the formula:

[(Q')(A)] or  $[(Q'')(B)_2(A)]$ 

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L P D

wherein

Q' is Pt(Y)2, where Y is an anion derived from a pharmaceutically acceptable acid, A is a diaminoacyl group having the formula

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where m is 0 or 1, n and p are 0-10, o is 1-10,

Q" is a non-platinum-group metal, either a main-group metal such as gallium, germanium, and tin, or a transition metal such as titanium, vanadium, iron, copper, cobalt, gold, nickel, cadmium and zinc,

B is a aralkylidene, heteroaralkylidene, cycloalkylidene or heterocycloalkylidene group containing oxygen anion or carboxylate anion at position 2 or 3, and pharmaceutically acceptable salts thereof and methods of use pertaining these compounds.



## EUROPEAN SEARCH REPORT

EP 89 11 8460

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